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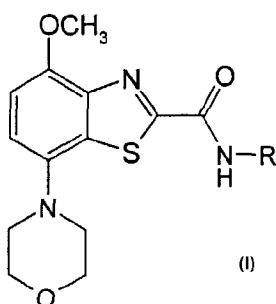
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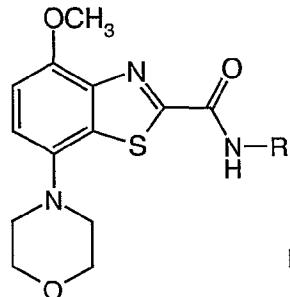


(57) **Abstract:** The invention relates to compounds of the general formula (I) wherein, R is hydrogen, -(CH₂)_n-phenyl, optionally substituted by halogen, lower alkyl, lower alkoxy, trifluoromethyl or -N(R')₂C(O)-lower alkyl, -(CH₂)_n-pyridinyl, optionally substituted by lower alkyl, -(CH₂)_n-C₃₋₆-cycloalkyl, optionally substituted by hydroxy, -(CH₂)_n-N(R')₂-C₃₋₆-cycloalkyl, -(CH₂)_n-benzo[1,3]-dioxolyl, -(CR')_n-thiophenyl, optionally substituted by lower alkyl, -(CR')_n-thiazolyl, optionally substituted by lower alkyl, -(CH₂)_n-C(O)-thiophenyl, optionally substituted by halogen, -(CH₂)_n-furanyl, optionally substituted by lower alkyl, -(CH₂)_n-C(O)-(CH₂)_n-thiophenyl, -(CHR')_n-benzofuran-2-yl, -(CH₂)_n-benzo[b]thiophenyl, optionally substituted by lower alkyl, -(CH₂)_n-N(R')₂-C(O)-phenyl, optionally substituted by halogen or lower alkoxy, -(CII₂)_n-C(O)-phenyl, optionally substituted by lower alkoxy, -(CII₂)_n-C(O)-2,3-dihydro-benzo [1,4]dioxin-6-yl, -(CH₂)_n-N(R')₂-C(O)-pyridinyl, -(CH₂)_n-tetrahydrofuranyl, -CH₂-bi-phenyl, -CH(phenyl)-pyridinyl, -(CH₂)_n-1-oxo-1,3-dihydro-isoindol-2-yl, -(CH₂)_n-1,3-dioxo-1,3-dihydro-isoindol-2-yl, -(CH₂)_n-CH(phenyl)-tetrahydropyranyl, -(CH₂)_n-1-oxo-1,2,3,4-tetrahydro-isoquinolin-3-yl or (CH₂)_n-S-[1,3,4]thiazol-2-yl, optionally substituted by amino; R' is hydrogen or lower alkyl, independently from each other in case R' is hydrogen or lower alkyl, independently from each other in case R' is hydrogen or lower alkyl, and pharmaceutically acceptable acid addition salts thereof. The compounds of formula I have a good affinity to the A_{2A} receptor and they are therefore useful for the treatment of diseases, related to this receptor.

Case 21015

Benzothiazole derivatives

The present invention relates to compounds of the general formula



wherein

R is

- hydrogen,
- $-(CH_2)_n$ -phenyl, optionally substituted by halogen, lower alkyl, lower alkoxy, trifluoromethyl or $-N(R')-C(O)$ -lower alkyl,
- $-(CH_2)_n$ -pyridinyl, optionally substituted by lower alkyl,
- $-(CH_2)_n-C_{3-6}$ -cycloalkyl, optionally substituted by hydroxy,
- $-(CH_2)_n-N(R')-C_{3-6}$ -cycloalkyl, optionally substituted by hydroxy,
- $-(CH_2)_n$ -benzo[1,3]-dioxolyl,
- $-(CR')_n$ -thiophenyl, optionally substituted by lower alkyl,
- $-(CR')_n$ -thiazolyl, optionally substituted by lower alkyl,
- $-(CH_2)_n-C(O)$ -thiophenyl, optionally substituted by halogen,
- $-(CH_2)_n$ -furanyl, optionally substituted by lower alkyl,
- $-(CH_2)_n-C(O)-(CH_2)_n$ -thiophenyl,
- $-(CHR')_n$ -benzofuran-2-yl,
- $-(CH_2)_n$ -benzo[b]thiophenyl, optionally substituted by lower alkyl,
- $-(CH_2)_n-N(R')-C(O)$ -phenyl, optionally substituted by halogen or lower alkoxy,
- $-(CH_2)_n-C(O)$ -phenyl, optionally substituted by lower alkoxy,
- $-(CH_2)_n-C(O)-2,3$ -dihydro-benzo[1,4]dioxin-6-yl,
- $-(CH_2)_n-N(R')-C(O)$ -pyridinyl,
- $-(CH_2)_n$ -tetrahydrofuranyl,
- $-CH$ -bi-phenyl,

-CH(phenyl)-pyridinyl,
-(CH₂)_n-1-oxo-1,3-dihydro-isoindol-2-yl,
-(CH₂)_n-1,3-dioxo-1,3-dihydro-isoindol-2-yl,
-(CH₂)_n-CH(phenyl)-tetrahydropyranyl,
5 -(CH₂)_n-1-oxo-1,2,3,4-tetrahydro-isoquinolin-3-yl or
-(CH₂)_n-S-[1,3,4]thiazol-2-yl, optionally substituted by amino;
R' is hydrogen or lower alkyl, independently from each other in case R'₂; and

n is 0, 1, 2, 3 or 4

and to pharmaceutically acceptable acid addition salts thereof.

10 It has surprisingly been found that the compounds of general formula I are adenosine receptor ligands. Specifically, the compounds of the present invention have a good affinity to the A_{2A}-receptor and a high selectivity to the A₁- and A₃ receptors.

15 Adenosine modulates a wide range of physiological functions by interacting with specific cell surface receptors. The potential of adenosine receptors as drug targets was first reviewed in 1982. Adenosine is related both structurally and metabolically to the bioactive nucleotides adenosine triphosphate (ATP), adenosine diphosphate (ADP), adenosine 20 monophosphate (AMP) and cyclic adenosine monophosphate (cAMP); to the biochemical methylating agent S-adenosyl-L-methione (SAM); and structurally to the coenzymes NAD, FAD and coenzym A; and to RNA. Together adenosine and these related compounds are important in the regulation of many aspects of cellular metabolism and in the modulation 25 of different central nervous system activities.

25 The receptors for adenosine have been classified as A₁, A_{2A}, A_{2B} and A₃ receptors, belonging to the family of G protein-coupled receptors. Activation of adenosine receptors by adenosine initiates signal transduction mechanism. These mechanisms are dependent on the receptor associated G protein. Each of the adenosine receptor subtypes has been classically characterised by the adenylyl cyclase effector system, which utilises cAMP as a second messenger. The A₁ and A₃ receptors, coupled with G_i proteins inhibit adenylyl cyclase, leading to a decrease in cellular cAMP levels, while A_{2A} and A_{2B} receptors couple to G_s proteins and activate adenylyl cyclase, leading to an increase in cellular cAMP levels. It 30 is known that the A₁ receptor system include the activation of phospholipase C and modulation of both potassium and calcium ion channels. The A₃ subtype, in addition to its association with adenylyl cyclase, also stimulates phospholipase C and so activates calcium ion channels.

The A₁ receptor (326-328 amino acids) was cloned from various species (canine, human, rat, dog, chick, bovine, guinea-pig) with 90-95 % sequence identify among the mammalian species. The A_{2A} receptor (409-412 amino acids) was cloned from canine, rat, human, guinea pig and mouse. The A_{2B} receptor (332 amino acids) was cloned from 5 human and mouse with 45 % homology of human A_{2B} with human A₁ and A_{2A} receptors. The A₃ receptor (317-320 amino acids) was cloned from human, rat, dog, rabbit and sheep.

The A₁ and A_{2A} receptor subtypes are proposed to play complementary roles in adenosine's regulation of the energy supply. Adenosine, which is a metabolic product of ATP, diffuses from the cell and acts locally to activate adenosine receptors to decrease the 10 oxygen demand (A₁) or increase the oxygen supply (A_{2A}) and so reinstate the balance of energy supply: demand within the tissue. The actions of both subtyps is to increase the amount of available oxygen to tissue and to protect cells against damage caused by a short term imbalance of oxygen. One of the important functions of endogenous adenosine is preventing damage during traumas such as hypoxia, ischaemia, hypotension and seizure 15 activity.

Furthermore, it is known that the binding of the adenosine receptor agonist to mast cells expressing the rat A₃ receptor resulted in increased inositol triphosphate and intracellular calcium concentrations, which potentiated antigen induced secretion of inflammatory mediators. Therefore, the A₃ receptor plays a role in mediating asthmatic 20 attacks and other allergic responses.

Adenosine is a neuromodulator, able to modulate many aspects of physiological brain function. Endogenous adenosine, a central link between energy metabolism and neuronal activity, varies according to behavioural state and (patho)physiological conditions. Under conditions of increased demand and decreased availability of energy 25 (such as hypoxia, hypoglycemia, and/or excessive neuronal activity), adenosine provides a powerful protective feedback mechanism. Interacting with adenosine receptors represents a promising target for therapeutic intervention in a number of neurological and psychiatric diseases such as epilepsy, sleep, movement disorders (Parkinson or Huntington's disease), Alzheimer's disease, depression, schizophrenia, or addiction. An increase in 30 neurotransmitter release follows traumas such as hypoxia, ischaemia and seizures. These neurotransmitters are ultimately responsible for neural degeneration and neural death, which causes brain damage or death of the individual. The adenosine A₁ agonists which mimic the central inhibitory effects of adenosine may therefore be useful as neuroprotective agents. Adenosine has been proposed as an endogenous anticonvulsant 35 agent, inhibiting glutamate release from excitatory neurons and inhibiting neuronal firing. Adenosine agonists therefore may be used as antiepileptic agents. Adenosine antagonists stimulate the activity of the CNS and have proven to be effective as cognition enhancers.

Selective A_{2a} antagonists have therapeutic potential in the treatment of various forms of dementia, for example in Alzheimer's disease, and of neurodegenerative disorders, e.g. stroke. Adenosine A_{2a} receptor antagonists modulate the activity of striatal GABAergic neurons and regulate smooth and well-coordinated movements, thus offering a potential 5 therapy for Parkinsonian symptoms. Adenosine is also implicated in a number of physiological processes involved in sedation, hypnosis, schizophrenia, anxiety, pain, respiration, depression, and drug addiction (amphetamine, cocaine, opioids, ethanol, nicotine, cannabinoids). Drugs acting at adenosine receptors therefore have therapeutic 10 potential as sedatives, muscle relaxants, antipsychotics, anxiolytics, analgesics, respiratory stimulants, antidepressants, and to treat drug abuse. They may also be used in the treatment of ADHD (attention deficit hyper-activity disorder).

An important role for adenosine in the cardiovascular system is as a cardioprotective agent. Levels of endogenous adenosine increase in response to ischaemia and hypoxia, and 15 protect cardiac tissue during and after trauma (preconditioning). By acting at the A₁ receptor, adenosine A₁ agonists may protect against the injury caused by myocardial ischemia and reperfusion. The modulating influence of A_{2a} receptors on adrenergic 20 function may have implications for a variety of disorders such as coronary artery disease and heart failure. A_{2a} antagonists may be of therapeutic benefit in situations in which an enhanced antiadrenergic response is desirable, such as during acute myocardial ischemia. Selective antagonists at A_{2a} receptors may also enhance the effectiveness of adenosine in terminating supraventricular arrhythmias.

Adenosine modulates many aspects of renal function, including renin release, glomerular filtration rate and renal blood flow. Compounds which antagonise the renal 25 affects of adenosine have potential as renal protective agents. Furthermore, adenosine A₃ and/or A_{2B} antagonists may be useful in the treatment of asthma and other allergic responses or and in the treatment of diabetes mellitus and obesity.

Numerous documents describe the current knowledge on adenosine receptors, for example the following publications:

Bioorganic & Medicinal Chemistry, 6, (1998), 619-641,
30 Bioorganic & Medicinal Chemistry, 6, (1998), 707-719,
J. Med. Chem., (1998), 41, 2835-2845,
J. Med. Chem., (1998), 41, 3186-3201,
J. Med. Chem., (1998), 41, 2126-2133,
J. Med. Chem., (1999), 42, 706-721,
35 J. Med. Chem., (1996), 39, 1164-1171,
Arch. Pharm. Med. Chem., 332, 39-41, (1999),

Am. J. Physiol., 276, H1113-1116, (1999) or
Naunyn Schmied, Arch. Pharmacol. 362, 375-381, (2000).

Objects of the present invention are the compounds of formula I per se, the use of compounds of formula I and their pharmaceutically acceptable salts for the manufacture of 5 medicaments for the treatment of diseases, related to the adenosine A₂ receptor, their manufacture, medicaments based on a compound in accordance with the invention and their production as well as the use of compounds of formula I in the control or prevention of illnesses based on the modulation of the adenosine system, such as Alzheimer's disease, 10 Parkinson's disease, Huntington's disease, neuroprotection, schizophrenia, anxiety, pain, respiration deficits, depression, drug addiction, such as amphetamine, cocaine, opioids, ethanol, nicotine, cannabinoids, or against asthma, allergic responses, hypoxia, ischaemia, 15 seizure and substance abuse. Furthermore, compounds of the present invention may be useful as sedatives, muscle relaxants, antipsychotics, antiepileptics, anticonvulsants and cardioprotective agents for disorders such as coronary artery disease and heart failure. The most preferred indications in accordance with the present invention are those, which base 20 on the A_{2A} receptor antagonistic activity and which include disorders of the central nervous system, for example the treatment or prevention of Alzheimer's disease, certain depressive disorders, drug addiction, neuroprotection and Parkinson's disease as well as ADHD.

As used herein, the term "lower alkyl" denotes a saturated straight- or branched- 25 chain alkyl group containing from 1 to 6 carbon atoms, for example, methyl, ethyl, propyl, isopropyl, n-butyl, i-butyl, 2-butyl, t-butyl and the like. Preferred lower alkyl groups are groups with 1 - 4 carbon atoms.

The term "halogen" denotes chlorine, iodine, fluorine and bromine.

The term "lower alkoxy" denotes a group wherein the alkyl residues is as defined 25 above, and which is attached via an oxygen atom.

The term "pharmaceutically acceptable acid addition salts" embraces salts with inorganic and organic acids, such as hydrochloric acid, nitric acid, sulfuric acid, phosphoric acid, citric acid, formic acid, fumaric acid, maleic acid, acetic acid, succinic acid, tartaric acid, methane-sulfonic acid, p-toluenesulfonic acid and the like.

30 Preferred compounds of the present application are compounds of formula I, wherein R is hydrogen, for example the following compound:
4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid amide.

Further preferred are compounds of formula I, wherein R is $-(CH_2)_n$ -phenyl, optionally substituted by halogen, lower alkoxy or lower alkyl, for example the following compounds:

- 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid phenethyl-amide,
- 5 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid 3-chloro-benzylamide,
- 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid 2-chloro-benzylamide,
- 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid 2-methoxy-benzylamide,
- 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(2-methoxy-phenyl)-ethyl]-amide,
- 10 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(3-fluoro-phenyl)-ethyl]-amide,
- 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(4-fluoro-phenyl)-ethyl]-amide,
- 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(4-chloro-phenyl)-ethyl]-amide,
- 15 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(2-chloro-phenyl)-ethyl]-amide,
- 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(3-methoxy-phenyl)-ethyl]-amide,
- 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(3-chloro-phenyl)-ethyl]-amide or
- 20 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-m-tolyl-ethyl)-amide.

Further preferred are compounds, wherein R is $-(CH_2)_n$ -pyridinyl, for example the following compounds:

- 25 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid pyridin-3-ylamide,
- 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (pyridin-2-ylmethyl)-amide or
- 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-pyridin-3-yl-ethyl)-amide.

- 30 Further preferred are compounds, wherein R is $-(CHR')_n$ -thiophenyl or $-(CH_2)_n-C(O)-$ thiophenyl, optionally substituted by halogen, for example the following compounds:

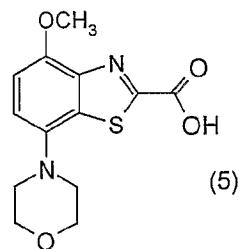
- 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-thiophen-2-yl-ethyl)-amide,
- 35 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-thiophen-3-yl-ethyl)-

amide,
 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(3-chloro-thiophen-2-yl)-2-oxo-ethyl]-amide or
 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (1-methyl-2-thiophen-2-yl-
 5 ethyl)-amide.

A preferred group of compounds is further those, wherein R is $-(CH_2)_n-thiazolyl$,
 optionally substituted by lower alkyl, for example the following compound:
 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [1-(4-methyl-thiazol-2-yl)-
 ethyl]-amide.

10 The present compounds of formula I and their pharmaceutically acceptable salts can
 be prepared by methods known in the art, for example, by processes described below,
 which processes comprise

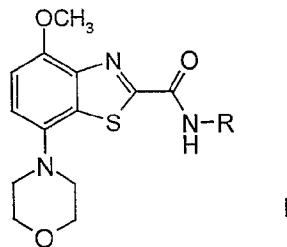
a) reacting a compound of formula



15 with a compound of formula

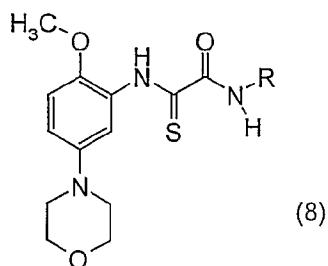


to a compound of formula

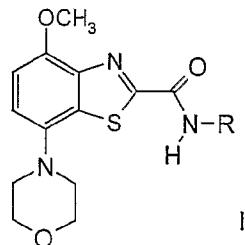


wherein R is as defined above, or

20 b) cyclising a compound of formula



to a compound of formula



wherein R is as described above, and

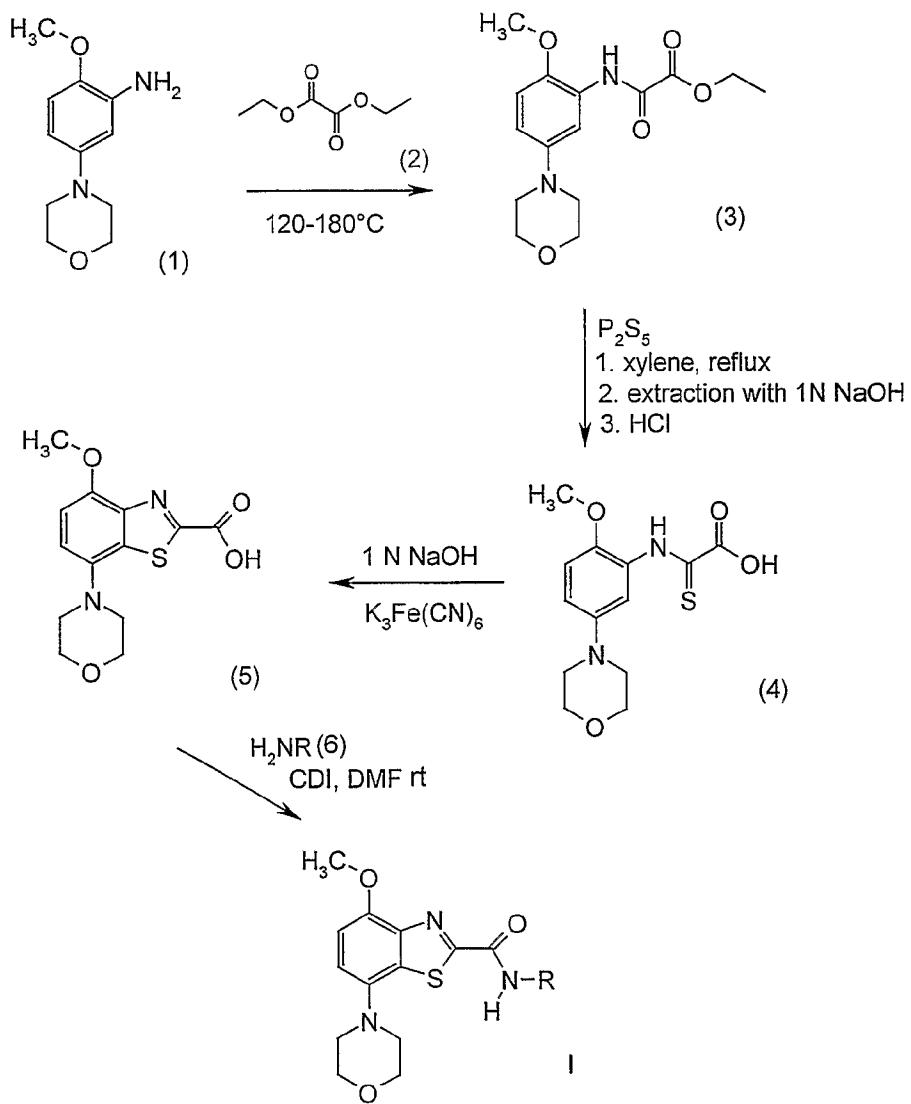
5 if desired, converting the compounds obtained into pharmaceutically acceptable acid addition salts.

The compounds of formula I may be prepared in accordance with process variants a) and b) and with the following schemes 1 and 2. The preparation of 84 Examples is further described in more detail.

10

15

Scheme 1



wherein R is as described above and CDI is 1,1'-carbonyl-diimidazole.

The preparation of the starting compound of formula (1) has been described in
5 WO 01/97786.

In accordance with scheme 1, the compounds of formula I are prepared as follows:

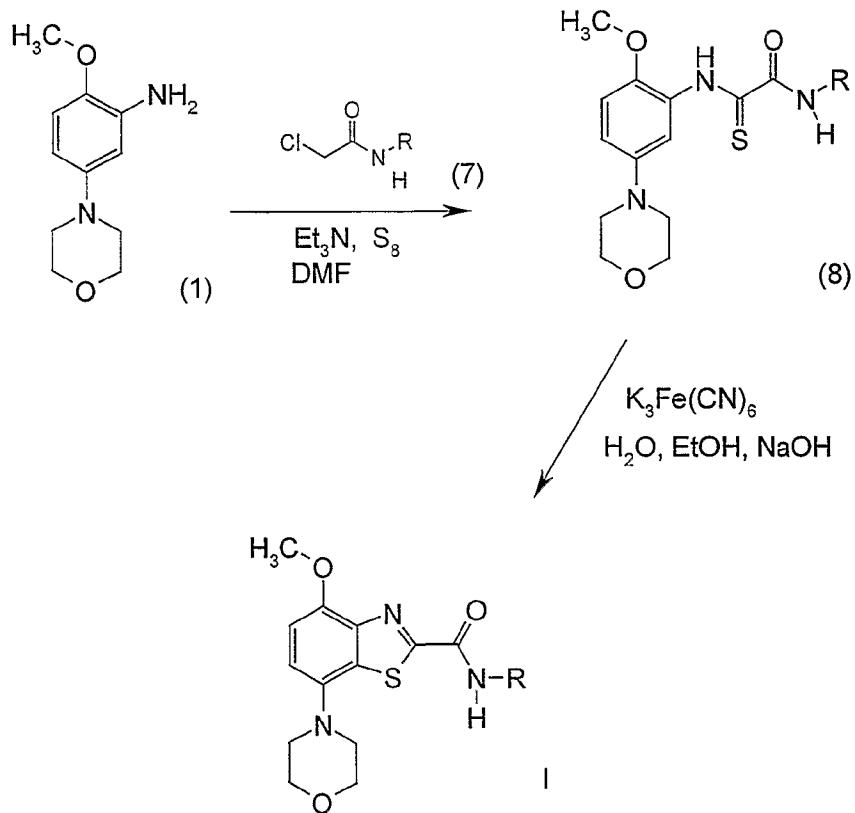
Diethyl oxalate (2) is heated to about 120 °C. 2-Methoxy-5-morpholin-4-yl-phenylamine (1) is added very cautiously in small quantities and the mixture is heated for 90 minutes at about 180 °C. After cooling to room temperature and filtration n-hexane is added. The 10 resulting precipitate is collected by filtration. After washing with hexane and drying N-(2-methoxy-5-morpholin-4-yl-phenyl)-oxalamic acid ethyl ester (3) is obtained. Then, to the obtained compound of formula (3) in boiling xylene is added phosphorus pentasulfide in

small portions over a period of about 30 minutes. The mixture is refluxed for about 5 hours, cooled to room temperature and filtered. The solution is extracted with 1N NaOH. The aqueous phase is washed with toluene, filtered, and treated at 0-5 °C with concentrated hydrochloric acid until pH 1 was reached. Filtration of the precipitate yielded (2-methoxy-5-morpholin-4-yl-phenylamino)-thioxo-acetic acid (4).

A solution of (2-methoxy-5-morpholin-4-yl-phenylamino)-thioxo-acetic acid (4) in 1N NaOH is added to a solution of potassium ferricyanide in water at a rate that the temperature does not exceed 10 °C. The mixture is stirred for 3 hours at 10 °C and concentrated hydrochloric acid is added until pH 1 was reached. Filtration of the 10 precipitate and drying yielded 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (5). A suspension of the compound of formula (5) and 1.1'-carbonyl-diimidazole in dimethylformamide is stirred at room temperature for about one hour. A compound of formula (6), for example benzylamine, is added, stirring is continued and after about 20 hours water is added. After extraction with ethyl acetate and chromatography on silicagel 15 with dichloromethane/ethylacetate is yielded a compound of formula I.

According to scheme 1, examples 3 to example 84 have been prepared.

Scheme 2



wherein R is as defined above.

In accordance with scheme 2 the corresponding chloroacetamide of formula (7) and sulfur in dimethylformamide are treated with triethylamine, and the mixture is stirred for about 15 hours at room temperature. Then 2-methoxy-5-morpholin-4-yl-phenylamine (1) and n-propanol are added and stirring at room temperature is continued for 6 hours. The 5 mixture is refluxed for two days. The precipitated crystals are filtered off and washed with n-propanol to yield a compound of formula (8).

A suspension of the compound of formula (8) in 1N aqueous sodiumhydroxide is added to a solution of potassium ferricyanide in water. The mixture is stirred at 50 °C for about 30 minutes and then at room temperature overnight. The precipitate is separated by filtration, 10 dissolved in dichloromethane and purified by column chromatography on silicagel with ethylacetate/hexane to yield a compound of formula I.

According to scheme 2, examples 1 and 2 have been prepared.

Isolation and purification of the compounds

Isolation and purification of the compounds and intermediates described herein can be effected, if desired, by any suitable separation or purification procedure such as, for 15 example, filtration, extraction, crystallization, column chromatography, thin-layer chromatography, thick-layer chromatography, preparative low or high-pressure liquid chromatography or a combination of these procedures. Specific illustrations of suitable separation and isolation procedures can be had by reference to the Preparations and 20 Examples herein below. However, other equivalent separation or isolation procedures could, of course, also be used.

Salts of compounds of formula I

The compounds of formula I may be basic, for example in cases where the residue R contains a basic group such as an aliphatic or aromatic amine moiety. In such cases the 25 compounds of Formula I may be converted to a corresponding acid addition salt.

The conversion is accomplished by treatment with at least a stoichiometric amount of an appropriate acid, such as hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid and the like, and organic acids such as acetic acid, propionic acid, glycolic acid, pyruvic acid, oxalic acid, malic acid, malonic acid, succinic acid, maleic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, cinnamic acid, mandelic acid, methanesulfonic acid, ethanesulfonic acid, p-toluenesulfonic acid, salicylic acid and the like. Typically, the free base is dissolved in an inert organic solvent such as diethyl ether, ethyl acetate, chloroform, ethanol or methanol and the like, and the acid added in a similar solvent. The 30

temperature is maintained between 0 °C and 50 °C. The resulting salt precipitates spontaneously or may be brought out of solution with a less polar solvent.

The acid addition salts of the basic compounds of Formula I may be converted to the corresponding free bases by treatment with at least a stoichiometric equivalent of a suitable 5 base such as sodium or potassium hydroxide, potassium carbonate, sodium bicarbonate, ammonia, and the like.

The compounds of formula I and their pharmaceutically usable addition salts possess valuable pharmacological properties. Specifically, it has been found that the compounds of the present invention are adenosine receptor ligands and possess a high affinity towards the 10 adenosine A_{2A} receptor and a good selectivity towards A₁ and A₃ receptors.

The compounds were investigated in accordance with the tests given hereinafter.

Human adenosine A₁ receptor

The human adenosine A₁ receptor was recombinantly expressed in chinese hamster ovary (CHO) cells using the semliki forest virus expression system. Cells were harvested, 15 washed twice by centrifugation, homogenised and again washed by centrifugation. The final washed membrane pellet was suspended in a Tris (50 mM) buffer containing 120 mM NaCl, 5 mM KCl, 2 mM CaCl₂ and 10 mM MgCl₂ (pH 7.4) (buffer A). The [³H]-DPCPX (([propyl-³H]8-cyclopentyl-1,3-dipropoxanthine); 0.6 nM) binding assay was carried out in 96-well plates in the presence of 2.5 µg of membrane protein, 0.5 mg of Ysi-poly-l-lysine 20 SPA beads and 0.1 U adenosine deaminase in a final volume of 200 µl of buffer A. Non-specific binding was defined using xanthine amine congener (XAC; 2 µM). Compounds were tested at 10 concentrations from 10 µM - 0.3 nM. All assays were conducted in duplicate and repeated at least two times. Assay plates were incubated for 1 hour at room temperature before centrifugation and then bound ligand determined using a Packard 25 Topcount scintillation counter. IC₅₀ values were calculated using a non-linear curve fitting program and Ki values calculated using the Cheng-Prussoff equation.

Human adenosine A_{2A} receptor

The human adenosine A_{2A} receptor was recombinantly expressed in chinese hamster ovary (CHO) cells using the semliki forest virus expression system. Cells were harvested, 30 washed twice by centrifugation, homogenised and again washed by centrifugation. The final washed membrane pellet was suspended in a Tris (50 mM) buffer containing 120 mM NaCl, 5 mM KCl, 2 mM CaCl₂ and 10 mM MgCl₂ (pH 7.4) (buffer A). The [³H]-SCH-58261 (Dionisotti et al., 1997, Br J Pharmacol 121, 353; 1nM) binding assay was carried out in 96-well plates in the presence of 2.5 µg of membrane protein, 0.5 mg of Ysi-poly-l-lysine

SPA beads and 0.1 U adenosine deaminase in a final volume of 200 μ l of buffer A. Non-specific binding was defined using xanthine amine congener (XAC; 2 μ M). Compounds were tested at 10 concentrations from 10 μ M - 0.3 nM. All assays were conducted in duplicate and repeated at least two times. Assay plates were incubated for 1 hour at room temperature before centrifugation and then bound ligand determined using a Packard Topcount scintillation counter. IC₅₀ values were calculated using a non-linear curve fitting program and Ki values calculated using the Cheng-Prusoff equation.

It has been shown that compounds of formula I have a good affinity to the A_{2A} receptor and a high selectivity toward the A₁. The preferred compounds show a pKi > 7.5.

Example No.	hA ₁ (pKi)	hA ₂ (pKi)
1	5.4	7.6
6	5.9	7.7
9	5.4	7.8
13	5.2	7.5
15	5.6	7.7
16	5.4	7.5
22	5.9	8.4
25	5.1	7.6
49	5.8	7.5
54	5.9	8.2
55	5.2	7.6
57	5.8	7.7
59	5.2	7.5
62	5.7	7.5
63	6.0	7.7

64	5.9	7.6
65	6.3	7.6
71	5.9	7.6
73	5.8	7.5
77	6.3	8.3
81	6.7	8.9

The compounds of formula I and the pharmaceutically acceptable salts of the compounds of formula I can be used as medicaments, e.g. in the form of pharmaceutical preparations. The pharmaceutical preparations can be administered orally, e.g. in the form of tablets, coated tablets, dragées, hard and soft gelatine capsules, solutions, emulsions or 5 suspensions. The administration can, however, also be effected rectally, e.g. in the form of suppositories, parenterally, e.g. in the form of injection solutions.

The compounds of formula I can be processed with pharmaceutically inert, inorganic or organic carriers for the production of pharmaceutical preparations. Lactose, corn starch or derivatives thereof, talc, stearic acids or its salts and the like can be used, for example, as 10 such carriers for tablets, coated tablets, dragées and hard gelatine capsules. Suitable carriers for soft gelatine capsules are, for example, vegetable oils, waxes, fats, semi-solid and liquid polyols and the like. Depending on the nature of the active substance no carriers are, however, usually required in the case of soft gelatine capsules. Suitable carriers for the 15 production of solutions and syrups are, for example, water, polyols, glycerol, vegetable oil and the like. Suitable carriers for suppositories are, for example, natural or hardened oils, waxes, fats, semi-liquid or liquid polyols and the like.

The pharmaceutical preparations can, moreover, contain preservatives, solubilizers, stabilizers, wetting agents, emulsifiers, sweeteners, colorants, flavorants, salts for varying 20 the osmotic pressure, buffers, masking agents or antioxidants. They can also contain still other therapeutically valuable substances.

Medicaments containing a compound of formula I or a pharmaceutically acceptable salt thereof and a therapeutically inert carrier are also an object of the present invention, as is a process for their production, which comprises bringing one or more compounds of formula I and/or pharmaceutically acceptable acid addition salts and, if desired, one or

more other therapeutically valuable substances into a galenical administration form together with one or more therapeutically inert carriers.

In accordance with the invention compounds of formula I as well as their pharmaceutically acceptable salts are useful in the control or prevention of illnesses based 5 on the adenosine receptor antagonistic activity, such as Alzheimer's disease, Parkinson's disease, neuroprotection, schizophrenia, anxiety, pain, respiration deficits, depression, asthma, allergic responses, hypoxia, ischaemia, seizure and substance abuse. Furthermore, compounds of the present invention may be useful as sedatives, muscle relaxants, 10 antipsychotics, antiepileptics, anticonvulsants and cardioprotective agents and for the production of corresponding medicaments.

The most preferred indications in accordance with the present invention are those, which include disorders of the central nervous system, for example the treatment or prevention of certain depressive disorders, neuroprotection and Parkinson's disease.

The dosage can vary within wide limits and will, of course, have to be adjusted to the 15 individual requirements in each particular case. In the case of oral administration the dosage for adults can vary from about 0.01 mg to about 1000 mg per day of a compound of general formula I or of the corresponding amount of a pharmaceutically acceptable salt thereof. The daily dosage may be administered as single dose or in divided doses and, in addition, the upper limit can also be exceeded when this is found to be indicated.

20

Tablet Formulation (Wet Granulation)

Item	Ingredients	mg/tablet			
		5 mg	25 mg	100 mg	500 mg
1.	Compound of formula I	5	25	100	500
2.	Lactose Anhydrous DTG	125	105	30	150
25	3.	Sta-Rx 1500	6	6	30
4.	Microcrystalline Cellulose	30	30	30	150
5.	Magnesium Stearate	1	1	1	1
	Total	167	167	167	831

Manufacturing Procedure

30 1. Mix items 1, 2, 3 and 4 and granulate with purified water.
 2. Dry the granules at 50°C.

3. Pass the granules through suitable milling equipment.
4. Add item 5 and mix for three minutes; compress on a suitable press.

Capsule Formulation

Item	Ingredients	<u>mg/capsule</u>			
		5 mg	25 mg	100 mg	500 mg
5	1. Compound of formula I	5	25	100	500
	2. Hydrous Lactose	159	123	148	---
	3. Corn Starch	25	35	40	70
	4. Talc	10	15	10	25
	10 5. Magnesium Stearate	1	2	2	5
Total		200	200	300	600

Manufacturing Procedure

1. Mix items 1, 2 and 3 in a suitable mixer for 30 minutes.
2. Add items 4 and 5 and mix for 3 minutes.
- 15 3. Fill into a suitable capsule.

The following preparation and examples illustrate the invention but are not intended to limit its scope.

Example 1

4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid amide

20 a) 2-(2-Methoxy-5-morpholin-4-yl-phenylamino)-2-thioxo-acetamide

173 mg (1.85 mmol) Chloroacetamide and 119 mg (3.70 mmol) sulfur in 2 ml dimethylformamide were treated with 772 μ l (5.55 mmol) triethylamine and the mixture was stirred for 15 hours at room temperature. Then 385 mg (1.85 mmol) 2-methoxy-5-morpholin-4-yl-phenylamine and 10 ml n-propanol were added and stirring at room temperature was continued for 6 hours. The mixture was refluxed for two days. The precipitated crystals were filtered off and washed with n-propanol to yield 320 mg (59 %) 2-(2-methoxy-5-morpholin-4-yl-phenylamino)-2-thioxo-acetamide as red-brown crystals. MS m/e (%): 296 ($M+H^+$, 100),

MA: $C_{13}H_{17}N_3O_3S$ (295.357) calc.: C 52.87 H 5.80 N 14.23 S 10.86

30 found: C 52.38 H 5.82 N 13.85 S 10.86

b) 4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid amide

A suspension of 220 mg (0.75 mmol) 2-(2-methoxy-5-morpholin-4-yl-phenylamino)-2-thioxo-acetamide in 2.88 ml 1N aqueous sodiumhydroxde was added to a solution of 813 mg (2.47 mmol) potassium ferricyanide in 2 ml water. The mixture was stirred at 50 °C for 5 30 minutes and then at room temperature overnight. The precipitate was separated by filtration, dissolved in dichloromethane and purified by column chromatography on silicagel with ethylacetate/hexane 1/1. 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid amide, 32 mg (15 %), was obtained as yellow crystals with mp.: 228-230 °C, MS m/e (%): 294 (M+H⁺, 100).

10

Example 2

4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (4-fluoro-phenyl)-amidea) N-(4-Fluoro-phenyl)-2-(2-methoxy-5-morpholin-4-yl-phenylamino)-2-thioxo-acetamide

919 mg (4.80 mmol) α -chloro-4-fluoroacetanilide and 308 mg (9.60 mmol) sulfur in 10 ml 15 dimethylformamide were treated with 2.01 ml (14.4 mmol) triethylamine and the mixture was stirred for 2 days at room temperature. Then 1.00 g (4.80 mmol) 2-methoxy-5-morpholin-4-yl-phenylamine in 5 ml dimethylformamide and 25 ml n-propanol were added and stirring at room temperature was continued for 6 hours. The mixture was refluxed for 6.5 hours. The precipitated crystals were filtered off and washed with water 20 and dried to yield 434 mg (24 %) N-(4-fluoro-phenyl)-2-(2-methoxy-5-morpholin-4-yl-phenylamino)-2-thioxo-acetamide as red powder with mp.: 206-208 °C, MS m/e (%): 390 (M+H⁺, 100).

b) 4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (4-fluoro-phenyl)-amide

25 A suspension of 100 mg (0.26 mmol) N-(4-fluoro-phenyl)-2-(2-methoxy-5-morpholin-4-yl-phenylamino)-2-thioxo-acetamide in 3.60 ml 1N aqueous sodiumhydroxde was added to a solution of 285 mg (0.87 mmol) potassium ferricyanide in 1 ml water. The mixture was stirred at 50 °C for two days and then extracted with dichloromethane. Purification by column chromatography on silicagel with ethylacetate/hexane 3/7 yielded 3.5 mg 4-30 methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (4-fluoro-phenyl)-amide as off-white crystals, MS m/e (%): 388 (M+H⁺, 100)

Example 3

4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid benzylamide

a) N-(2-Methoxy-5-morpholin-4-yl-phenyl)-oxalamic acid ethyl ester

139 ml (1015 mmol) Diethyl oxalate were heated to 120 °C. 30.3 g (145 mmol) 2-methoxy-5-morpholin-4-yl-phenylamine were added very cautiously in small quantities and the mixture was heated for 90 minutes at 180 °C. After cooling to room temperature and 5 filtration 1.5 l n-hexane were added. The resulting precipitate was collected by filtration. After washing with hexane and drying 34.4 g (77 %) N-(2-methoxy-5-morpholin-4-yl-phenyl)-oxalamic acid ethyl ester was obtained as greenish crystalline, mp.: 95-97 °C, MS m/e (%): 309 (M+H⁺, 100).

b) (2-Methoxy-5-morpholin-4-yl-phenylamino)-thioxo-acetic acid

10 To 33.9 g (110 mmol) N-(2-methoxy-5-morpholin-4-yl-phenyl)-oxalamic acid ethyl ester in 652 ml boiling xylene were added 8.80 g (40 mmol) phosphorus pentasulfide in small portions over a period of 30 minutes. The mixture was refluxed for 5 hours, cooled to room temperature and filtered. The solution was extracted 7 times with 100 ml 1N NaOH. The aqueous phase was washed twice with 100 ml toluene, filtered, and treated at 0-5 °C 15 with concentrated hydrochloric acid until pH 1 was reached. Filtration of the precipitate yielded 20.2 g (62 %) (2-methoxy-5-morpholin-4-yl-phenylamino)-thioxo-acetic acid as yellow crystalline, mp.: 156-158 °C, MS m/e (%): 297 (M+H⁺, 100).

c) 4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid

20 A solution of 10.5 g (35.4 mmol) (2-methoxy-5-morpholin-4-yl-phenylamino)-thioxo-acetic acid in 248 ml (248 mmol) 1N NaOH was added dropwise to a solution of 40.1 g (119 mmol) potassium ferricyanide in 119 ml water at a rate that the temperature did not exceed 10 °C. The mixture was stirred for 3 hours at 10 °C and concentrated hydrochloric acid was added until pH 1 was reached. Filtration of the precipitate and drying yielded 8.80 g (84 %) 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid as yellow crystalline 25 with mp.: 99-100 °C, MS m/e (%): 295 (M+H⁺, 100).

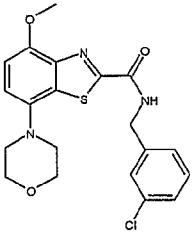
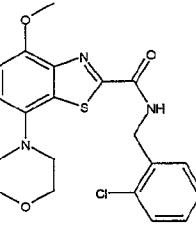
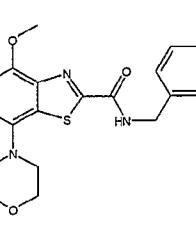
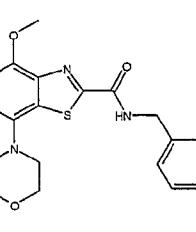
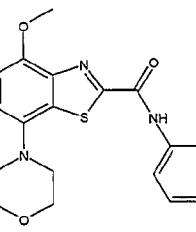
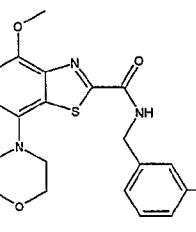
d) 4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid benzylamide

30 A suspension of 29.4 mg (0.10 mmol) 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid and 18.4 mg (0.11 mmol) 1,1'-carbonyl-diimidazole in 3 ml dimethylformamide was stirred at room temperature for one hour. 12.1 µl (0.11 mmol) benzylamine were added, stirring was continued and after 20 hours 15 ml water were added. Extraction with ethyl acetate and chromatography on silicagel with dichloromethane/ethylacetate 95/5 yielded 31.2 mg (41 %) of yellow 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid benzylamide with mp.: 156-158 °C, MS m/e (%): 384 (M+H⁺, 100).

According to example 3d derivatives have been synthesised. They are compiled in the following list comprising example 4 to example 81:

Exp. No.	Structure	Systematic name	m.p. °C	educt
4		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid phenylamide	190-191	Aniline
5		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid cyclohexylamide	146-147	Cyclohexyl-amine
6		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid phenethyl-amide	50-51	Phenylethyl-amine
7		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid 4-chloro-benzylamide	175-176	4-Chloro-benzylamine
8		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid cyclopentylamide	174-175	Cyclopentyl-amine

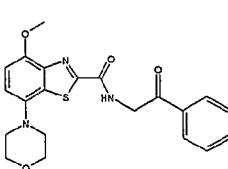
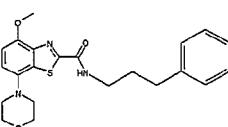
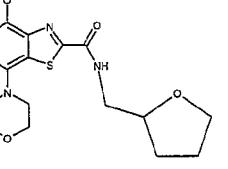
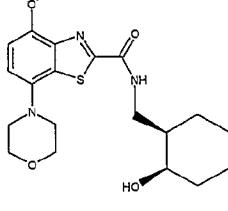
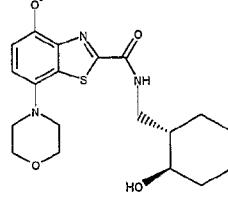
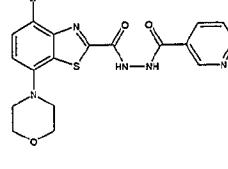
9		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid pyridin-3-ylamide	220-221	3-Amino-pyridin
10		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (4-methoxy-phenyl)-amide	148	p-Anisidin
11		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (3-methoxy-phenyl)-amide	160	m-Anisidin
12		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (2-methoxy-phenyl)-amide	191	o-Anisidin
13		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (pyridin-2-ylmethyl)-amide	181-182	2-Amino-methyl-pyridine
14		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (benzo[1,3]dioxol-5-ylmethyl)-amide	190-193	1,3-Benzo-dioxole-5-methyl-amine

15		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid 3-chloro-benzylamide	148-150	3-Chlorobenzylamine
16		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid 2-chloro-benzylamide	114-116	2-Chlorobenzylamine
17		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid 4-fluoro-benzylamide	155-156	4-Fluorobenzylamine
18		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (pyridin-3-ylmethyl)-amide	181-184	3-(Aminomethyl)-pyridine
19		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid pyridin-2-ylamide	177-181	2-Aminopyridine
20		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid 3-fluoro-benzylamide	152-153	3-Fluorobenzylamine

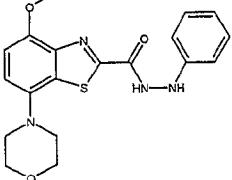
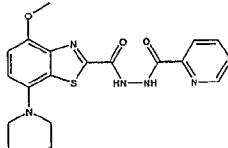
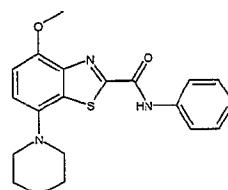
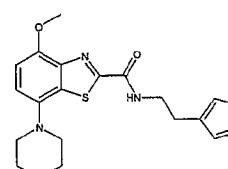
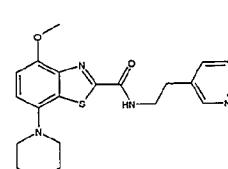
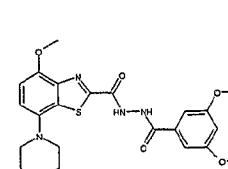
21		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid 2-fluoro-benzylamide	133-134	2-Fluorobenzylamine
22		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (2-thiophen-2-yl-ethyl)-amide	120-121	2-(2-Thienyl)-ethylamine
23		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (thiophen-2-ylmethyl)-amide	174-175	2-Thiophenmethylamine
24		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid cyclopropylmethyl-amide	152-153	(Amino-methyl)-cyclopropane
25		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid 2-methoxy-benzylamide	186-188	2-Methoxybenzylamine
26		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid 4-methoxy-benzylamide	164-168	4-Methoxybenzylamine

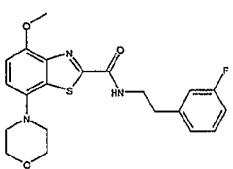
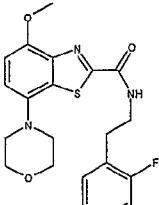
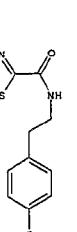
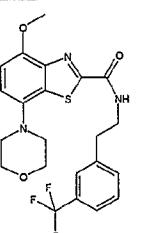
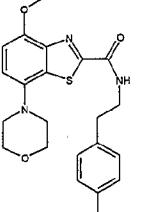
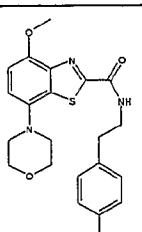
27		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid 3-methoxy-benzylamide	140-144	3-Methoxybenzylamine
28		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(4-chloro-benzoylamino)-ethyl]-amide	237-240	N-(2-Aminoethyl)-p-chlorobenzamide
29		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(1-oxo-1,3-dihydro-isoindol-2-yl)-ethyl]-amide	199-205	2-(2-Aminoethyl)-phthal-imidine
30		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (4-phenyl-tetrahydro-pyran-4-ylmethyl)-amide	144-148	4-Phenyl-4-methyl-amino-tetrahydro-pyran
31		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (pyridin-4-ylmethyl)-amide	207-209	4-Picolyamine
32		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [3-(1,3-dioxo-1,3-dihydro-isoindol-2-yl)-propyl]-amide	200-202	N-(3-Aminopropyl)-phthalimide

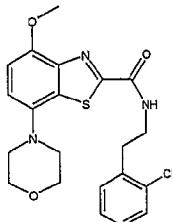
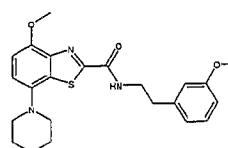
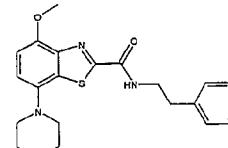
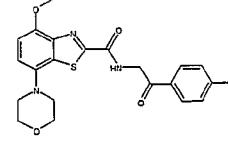
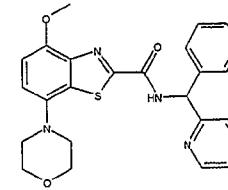
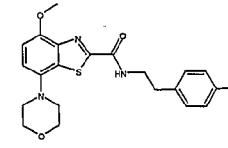
33		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [4-(1,3-dioxo-1,3-dihydro-isoindol-2-yl)-butyl]-amide	114-118	N-(4-Amino-butyl)-phthalimide
34		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(1,3-dioxo-1,3-dihydro-isoindol-2-yl)-ethyl]-amide	231-233	N-(2-Aminoethyl)-phthalimide
35		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid cyclohexylmethyl-amide	150-151	Cyclohexyl-methylamine
36		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (1-oxo-1,2,3,4-tetrahydro-isoquinolin-3-ylmethyl)-amide	152-155	3-(Amino-methyl)-3,4-dihydro-1(2H)-iso-quinolinone
37		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(2-methoxy-benzoylamino)-ethyl]-amide	180-183	N-(2-Aminoethyl)-o-anisamide
38		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-pyridin-2-yl-ethyl)-amide	184-185	2-(2-Pyridyl)-ethylamine

39		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-oxo-2-phenyl-ethyl)-amide	197-200	Aminoaceto-phenone
40		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (3-phenyl-propyl)-amide	147	3-Phenyl-propylamine
41		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (tetrahydro-furan-2-ylmethyl)-amide	171-172	Tetrahydro-furfuryl-amine
42		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-hydroxy-cyclohexylmethyl)-amide	178-180	cis-2-Amino-ethyl-1-cyclohexanol
43		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-hydroxy-cyclohexylmethyl)-amide	133-135	trans-2-Aminoethyl-1-cyclo-hexanol
44		Nicotinic acid N'-(4-methoxy-7-morpholin-4-yl-benzothiazole-2-carbonyl)-hydrazide	222-224	Nicotinic acid hydrazide

45		Isonicotinic acid N'-(4-methoxy-7-morpholin-4-yl-benzothiazole-2-carbonyl)-hydrazide	171-174	Isonicotinic acid hydrazide
46		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid N'-benzoyl-hydrazide	>260°C	Benz-hydrazide
47		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid benzhydryl-amide	166-168	alpha-Amino-diphenyl-methane
48		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-pyridin-4-yl-ethyl)-amide	185-188	4-(2-Aminoethyl)-pyridine
49		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(2-methoxy-phenyl)-ethyl]-amide	158-160	2-(2-Methoxy-phenyl)-ethylamine
50		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(3-acetylamino-phenyl)-ethyl]-amide	165-166	2-(3-Acetylamino-phenyl)-ethylamine

51		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid N'-phenyl-hydrazide	226-227	Phenyl-hydrazine
52		Pyridine-2-carboxylic acid N'-(4-methoxy-7-morpholin-4-ylbenzothiazole-2-carbonyl)-hydrazide	225-226	2-Picolinyl hydrazide
53		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid pyridin-4-ylamide	206-207	4-Amino-pyridine
54		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (2-thiophen-3-yl-ethyl)-amide	137-139	3-Thiophene-ethylamine
55		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (2-pyridin-3-yl-ethyl)-amide	160-163	3-(2-Aminoethyl)-pyridine
56		3,5-Dimethoxy-benzoic acid N'-(4-methoxy-7-morpholin-4-ylbenzothiazole-2-carbonyl)-hydrazide	146-148	3,5-Dimethoxy-benz-hydrazide

57		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(3-fluoro-phenyl)-ethyl]-amide	120-122	3-Fluorophenylethylamine
58		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(2-fluoro-phenyl)-ethyl]-amide	151-153	2-Fluorophenylethylamine
59		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(4-fluoro-phenyl)-ethyl]-amide	150-153	4-Fluorophenylethylamine
60		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(3-trifluoromethyl-phenyl)-ethyl]-amide	106-108	2-(3-Trifluoromethylphenyl)-ethylamine
61		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (2-p-tolyl-ethyl)-amide	136-138	4-Methylphenylethylamine
62		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(4-chloro-phenyl)-ethyl]-amide	153-154	2-(4-Chlorophenyl)-ethylamine

63		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(2-chlorophenyl)-ethyl]-amide	141-143	2-(2-Chlorophenyl)-ethylamine
64		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(3-methoxyphenyl)-ethyl]-amide	99-101	3-Methoxyphenylethylamine
65		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(3-chlorophenyl)-ethyl]-amide	109-111	3-Chlorophenylethylamine
66		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(4-methoxyphenyl)-2-oxoethyl]-amide	176-178	2-amino-4'-methoxyaceto-phenone
67		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (phenyl-pyridin-2-yl-methyl)-amide	219-221	Phenyl-(2-pyridyl)-methylamine
68		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(4-methoxyphenyl)-ethyl]-amide	69-71	4-Methoxyphenylethylamine

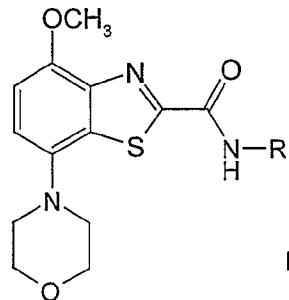
69		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (benzo[b]thiophen-3-ylmethyl)-amide	187-189	Benzo(b)thiophen-3-ylmethylamine
70		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (5-methyl-furan-2-ylmethyl)-amide	173-174	5-Methylfurylamine
71		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (2-m-tolyl-ethyl)-amide	119-121	3-Methylphenethylamine
72		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (2-hydroxy-cyclohexyl)-amide	110-115	trans-2-Aminocyclohexanol
73		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(3-chloro-thiophen-2-yl)-2-oxo-ethyl]-amide	125-127	2-(3-Chlorothien-2-yl)-2-oxo-1-ethanamine
74		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (2-o-tolyl-ethyl)-amide	129-131	2-Methylphenethylamine

75		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(2,3-dihydrobenzo[1,4]dioxin-6-yl)-2-oxoethyl]-amide	206-208	2-Amino-1-(2,3-dihydrobenzo[1,4]-dioxin-6-yl)-ethanone
76		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [2-(3-methyl-benzo[b]thiophen-2-yl)-2-oxoethyl]-amide	185-187	2-Amino-1-(3-methyl-benzo[b]-thiophen-2-yl)-ethanone
77		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [1-(4-methyl-thiazol-2-yl)-ethyl]-amide	175-176	alpha-Methyl-2-(4-methyl-thiazole)-methan-amine
78		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (2-oxo-2-thiophen-2-yl-ethyl)-amide	171-173	2-Amino-1-(2-thienyl)ethanone
79		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid (2-oxo-2-thiophen-3-yl-ethyl)-amide	229-232	2-Amino-1-(3-thienyl)ethanone
80		4-Methoxy-7-morpholin-4-ylbenzothiazole-2-carboxylic acid [1-(2-methyl-thiazol-4-yl)-ethyl]-amide	174-175	4-(1-Aminoethyl)-2-methyl-thiazole

81		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (1-methyl-2-thiophen-2-yl-ethyl)-amide	127-129	2-(2-Amino-propyl)-thiophene
82		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (6-methyl-pyridin-3-yl)-amide	145-147	5-Amino-2-methylpyridine
83		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(5-amino-[1,3,4]thiadiazol-2-ylsulfanyl)-ethyl]-amide	237-239	2Amino-5-[(2-aminoethyl)thio]-1,3,4-thiadiazole
84		4-Methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-benzofuran-2-yl-1-methyl-ethyl)-amide	144-146	2-(2-Aminopropyl)-benzofuran

Claims

1. Compounds of the general formula



wherein

5 R is
 hydrogen,
 $-(CH_2)_n$ -phenyl, optionally substituted by halogen, lower alkyl, lower alkoxy, trifluoromethyl or $-N(R')-C(O)$ -lower alkyl,
 $-(CH_2)_n$ -pyridinyl, optionally substituted by lower alkyl,

10 $-(CH_2)_n-C_{3-6}$ -cycloalkyl, optionally substituted by hydroxy,
 $-(CH_2)_n-N(R')-C_{3-6}$ -cycloalkyl, optionally substituted by hydroxy,
 $-(CH_2)_n$ -benzo[1,3]-dioxolyl,
 $-(CR')_n$ -thiophenyl, optionally substituted by lower alkyl,
 $-(CR')_n$ -thiazolyl, optionally substituted by lower alkyl,

15 $-(CH_2)_n-C(O)$ -thiophenyl, optionally substituted by halogen,
 $-(CH_2)_n$ -furanyl, optionally substituted by lower alkyl,
 $-(CH_2)_n-C(O)-(CH_2)_n$ -thiophenyl,
 $-(CHR')_n$ -benzofuran-2-yl,
 $-(CH_2)_n$ -benzo[b]thiophenyl, optionally substituted by lower alkyl,

20 $-(CH_2)_n-N(R')-C(O)$ -phenyl, optionally substituted by halogen or lower alkoxy,
 $-(CH_2)_n-C(O)$ -phenyl, optionally substituted by lower alkoxy,
 $-(CH_2)_n-C(O)-2,3$ -dihydro-benzo[1,4]dioxin-6-yl,
 $-(CH_2)_n-N(R')-C(O)$ -pyridinyl,
 $-(CH_2)_n$ -tetrahydrofuranyl,

25 $-CH$ -bi-phenyl,
 $-CH(phenyl)$ -pyridinyl,
 $-(CH_2)_n$ -1-oxo-1,3-dihydro-isoindol-2-yl,
 $-(CH_2)_n$ -1,3-dioxo-1,3-dihydro-isoindol-2-yl,

-(CH₂)_n-CH(phenyl)-tetrahydropyranyl,
-(CH₂)_n-1-oxo-1,2,3,4-tetrahydro-isoquinolin-3-yl or
-(CH₂)_n-S-[1,3,4]thiazol-2-yl, optionally substituted by amino;
R' is hydrogen or lower alkyl, independently from each other in case R'₂, and
5 n is 0, 1, 2, 3 or 4;
and pharmaceutically acceptable acid addition salts thereof.

2. Compounds of formula I in accordance with claim 1, wherein R is hydrogen.

3. A compound of formula I in accordance with claim 2, wherein the compound is 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid amide.

10 4. Compounds in accordance with claim 1, wherein R is -(CH₂)_n-phenyl, optionally substituted by halogen, lower alkoxy or lower alkyl.

5. Compounds in accordance with claim 4, wherein the compound is 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid phenethyl-amide,
4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid 3-chloro-benzylamide,
15 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid 2-chloro-benzylamide,
4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid 2-methoxy-benzylamide,
4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(2-methoxy-phenyl)-ethyl]-amide,
4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(3-fluoro-phenyl)-
20 ethyl]-amide,
4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(4-fluoro-phenyl)-ethyl]-amide,
4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(4-chloro-phenyl)-ethyl]-amide,
25 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(2-chloro-phenyl)-ethyl]-amide,
4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(3-methoxy-phenyl)-ethyl]-amide,
4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(3-chloro-phenyl)-
30 ethyl]-amide or
4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-m-tolyl-ethyl)-amide.

6. Compounds in accordance with claim 1, wherein R is $-(CH_2)_n$ -pyridinyl.

7. Compounds in accordance with claim 6, wherein the compound is
 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid pyridin-3-ylamide,
 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (pyridin-2-ylmethyl)-amide
 5 or
 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-pyridin-3-yl-ethyl)-amide.

8. Compounds in accordance with claim 1, wherein R is $-(CHR')_n$ -thiophenyl or
 $-(CH_2)_n-C(O)$ -thiophenyl, optionally substituted by halogen.

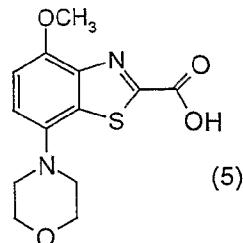
10 9. Compounds in accordance with claim 8, wherein the compound is
 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-thiophen-2-yl-ethyl)-amide,
 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (2-thiophen-3-yl-ethyl)-amide,
 15 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [2-(3-chloro-thiophen-2-yl)-2-oxo-ethyl]-amide or
 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid (1-methyl-2-thiophen-2-yl-ethyl)-amide.

20 10. Compounds in accordance with claim 1, wherein R is $-(CHR')_n$ -thiazolyl,
 optionally substituted by lower alkyl.

11. Compounds in accordance with claim 10, wherein the compound is
 4-methoxy-7-morpholin-4-yl-benzothiazole-2-carboxylic acid [1-(4-methyl-thiazol-2-yl)-ethyl]-amide.

25 12. A process for preparing a compound of formula I as defined in claims 1 - 11,
 which processes comprise

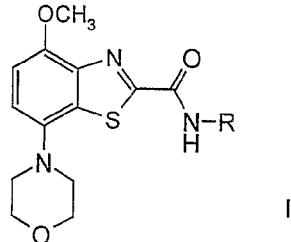
a) reacting a compound of formula



with a compound of formula

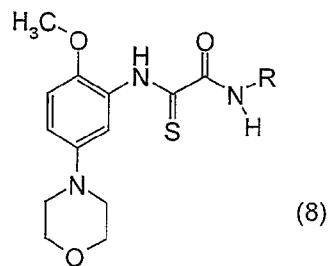


to a compound of formula

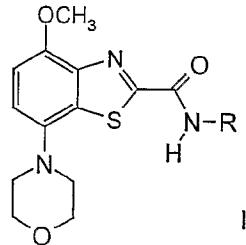


wherein R is as defined in claim 1, or

5 b) cyclising a compound of formula



to a compound of formula



wherein R is as described in claim 1, and

10 if desired, converting the compounds obtained into pharmaceutically acceptable acid addition salts.

13. A compound according to any one of claims 1 to 11, whenever prepared by a process as claimed in claim 14 or by an equivalent method.

14. A medicament containing one or more compounds as claimed in any one of claims 1 to 11 and pharmaceutically acceptable excipients.

15. A medicament according to claim 14 for the treatment of diseases related to the adenosine receptor.

16. The use of a compound in any one of claims 1 to 11 for the treatment of diseases.
17. The use of a compound in any one of claims 1 to 11 for the manufacture of corresponding medicaments for the treatment of diseases related to the adenosine A_{2A} receptor.
- 5 18. The invention as hereinbefore described.

INTERNATIONAL SEARCH REPORT

Int'l	Application No
PCT/EP 02/12907	

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7	A61K31/428	A61K31/429	A61K31/44	A61P25/16	A61P25/28
	C07D277/82	C07D417/12	C07D409/12	C07D477/12	

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, CHEM ABS Data, BEILSTEIN Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	<p>WO 01 97786 A (HOFFMANN LA ROCHE) 27 December 2001 (2001-12-27) claims 1,19</p> <p>-----</p>	1-17



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority, claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

& document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the International search report
22 January 2003	29/01/2003
Name and mailing address of the ISA	Authorized officer

European Patent Office, P.B. 5818 Patentlaan 2

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Gettins, M

INTERNATIONAL SEARCH REPORT

International application No.
PCT/EP 02/12907

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:

2. Claims Nos.: 18 because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210

3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this International application, as follows:

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 18

The scope of claim 18 "The invention as hereinbefore described" is either merely a repetition of claims 1-17 in which case it is superfluous or a fully unclear reference to the description (Art 6 PCT). In either case it has not been searched.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

Information on patent family members

Int'l Application No

PCT/EP 02/12907

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 0197786	A 27-12-2001	AU 8181701 A WO 0197786 A2 US 2002045615 A1	02-01-2002 27-12-2001 18-04-2002